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(54) Trocar wound closure device

Trokarwundverschlussvorrichtung

Dispositif pour obturer des plaies de trocart

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Description

This invention relates generally to devices for facilitating the suturing of trocar puncture wounds, and more particularly to a device that is insertable through a cannula for presenting a pair of needles loaded onto the ends of a suture, in position to close the trocar wound from within an insufflated or otherwise distended abdominal cavity.

Laparoscopic surgery commonly includes the creation of one or more trocar puncture wounds through the anterior abdominal wall for receiving surgical instruments. The structural strength of the abdominal wall is derived mostly from one or more layers of fascia (connective tissue sheets) running parallel to skin and between layers of muscle. The increasing complexity of surgical procedures performed laparoscopically has led to the development and use of larger diameter trocars. Trocars having outside diameters of up to 12 mm. are currently being used, and we anticipate that trocars of up to 20 mm or larger will be used in the future.

While small trocar puncture wounds will often heal satisfactorily without suturing, the wounds created by larger trocars may not. Unless closed properly, larger diameter trocar wounds may allow abdominal contents to herniate into a fascial defect.

Presently, surgeons attempt to suture trocar puncture wounds using conventional hand held needle drivers external to the patient. Because of the limited exposure and the potential for puncturing abdominal organs, conventional needle driving techniques typically place sutures only through the outer layers of fascia. These manual techniques are often further compromised by patient obesity and other factors.

Accordingly, there is a need for an improved method of closing trocar puncture wounds to minimize the risk of puncture site herniation or other undesirable side effects of known trocar wound closure techniques.

US-A-5047039 discloses an anastomosis apparatus with a pair of needles and a suture mounted at its distal end. DE-A-4137218 falls to be considered in Germany as a document which, although unpublished on the filing date, declares an earlier priority date. In its patent family is EP-A-542126.

Briefly stated, a trocar wound closure device according to this invention includes an elongated body and first and second retractable needle holders disposed at the distal end of the body. The needle holders are movable between a retracted position and an extended position. An actuator disposed at the proximal end of the body moves the needle holders from the retracted position to the extended position, so that the needle holders can be retracted to allow the device to be inserted through a trocar wound preferably through a cannula inserted into the wound, and extended to position the needles adjacent the wound, to allow the wound to be sutured.

The elongated body tube may be a tubular body, and the needle holders may be elongated cantilevered arms fixed at the proximal end, and free at the distal end.

The retractable needle holders are movable from a retracted position within the tubular body to an extended position outside the tubular body.

The needle holders may have tapered retracting edges for causing the needle holders to retract as the wound closure device is withdrawn from a trocar wound, especially through a cannula.

The actuator may comprise a plunger movable within the tubular body from an extended position to a depressed position, for engaging and moving the cantilevered arms outward as the plunger is depressed, and moving the needle holders from the retracted position to the extended position, respectively.

The needle holders may be needle plates slidably mounted in the elongated body, with needle holding edges oriented generally parallel to the body.

The body may include a tube for carrying a sutur within the body.

While the novel aspects of this invention are set forth with particularity in the appended claims, the invention itself, together with further objects and advantages thereof, may be more readily comprehended by reference to the following detailed description of a presently preferred embodiment of the invention, taken in conjunction with the following drawings, in which:

Figure 1 is a side elevational view of a trocar wound closure device in accordance with this invention;
Figure 2 is a side elevation of the plunger of the trocar wound closure device of Figure 1;
Figure 3 is a bottom plan view of the plunger of Figure 2;
Figure 4 is an enlarged segmental view of the distal end of the trocar wound closure device of Figure 1, shown partly in section, and loaded with a suture;
Figure 5 is a side elevation of the distal end of the trocar wound closure device shown in Figure 4;
Figure 6 is a bottom plan view of the distal end of the trocar wound closure device of Figure 4 showing the needle holders in the extended position; and
Figure 7 is a bottom plan view similar to Figure 6, but showing the needle holders in the retracted position with the suture and needles omitted for clarity;
Figure 8 is an enlarged view of the proximal end of the trocar wound closure device, shown partly in section, with the plunger fully depressed;
Figure 9 is an enlarged side elevational view of the proximal end of the trocar wound closure device sirnilar to that shown in Figure 8, but with the plunger extended;
Figure 10 is a simplified side elevational view of a cannula for use with a trocar wound closure device in accordance with this invention;
Figure 11 is a side elevational view, partly in section, showing the cannula of Figure 10 inserted through a trochar wound into a peritoneal cavity

with the trocar wound closure device of this invention in the process of being inserted into the cannula;

Figure 12 is a side elevational view, partly in section, showing the trocar wound closure device fully inserted into the cannula with the needle holders extended;

Figure 13 is a side elevational view, partly in section, showing the process of withdrawing the wound closure device and cannula to cause the needles to pierce the peritoneal wall;

Figure 14 is a fragmentary side elevational view showing the distal end of the trocar wound closure device as both sides of the the suture are pulled through the peritoneal wall;

Figure 15 is a fragmentary side elevational view showing the trocar wound closure device with the needle holders retracted in preparation for withdrawal from the trocar puncture wound site;

Figure 16 is a perspective view of an alternative embodiment of the distal end of a trocar wound closure device;

Figure 17 is a section thereof taken along lines 17-17 of Figure 16 showing the needle holders rotated to an extended position;

Figure 18 is a section taken along the same line showing the needle holders rotated to a retracted position; and

Figure 19 is a section taken along line 19-19 of Figure 16 showing the distal end of the alternative embodiment of the trocar wound closure device.

In a presently preferred embodiment of the invention as shown in Figure 1, the trocar wound closure device 10 includes an elongated tubular, preferably cylindrical, body 12 having a proximal end 14 inserted into a housing block 16, preferably formed from a moldable or machinable plastic, soft metal or the like, and a distal end 20. A plunger 22 having a handle 24 attached thereto at one end 26 is slidably received within the tubular body 12. The plunger 22 is shown in a side elevation in Figure 3, and a bottom plan view of the plunger is shown in Figure 2. The plunger 22 includes the handle 24 already described, and an elongated shaft 28 attached to the handle by conventional fasteners, such as a screw or rivet 30.

An actuating element 32 is attached to the distal end 34 of the elongated shaft 28. The actuating element 32 is preferably machined or molded from a low coefficient of friction plastic material such as delrin or the like. It is generally cylindrical in cross section, and has three longitudinally extending notches 40, 42, 44 formed therein, spaced radially around the periphery of the actuating element 32. A first shallow notch 40 and a second shallow notch 42 are adapted to engage first and second needle drivers as will be shown and described in more detail later. A deeper notch 44 receives a suture holder.

The distal end 20 of the trocar wound closure device 10 is shown in more detail in Figure 4, which is partly broken away and sectioned for convenience. First and second elongated slots 50 and 52 are formed in the distal end 20 of the tubular body 12, as can be seen also in Figures 5, 6, and 7. The slots 50 and 52 extend from the distal end 20 of the body 12 towards the proximal end 14 a distance sufficient to allow first and second needles 56 and 58 to be retracted laterally into the body through the slots 50 and 52.

The needles 56 and 58 are carried by needle drivers having first and second elongated resilient bars 64 and 66. The bars have first and second fin shaped needle carriers 70 and 72 attached at the distal ends of the bars 64 and 66. Each of the needle holders has a long side 74, 76 attached to the respective distal end of its bar, and a remote shorter side 78, 80 having a needle carrying groove 82, 84 formed therein. The top sides 88, 90 of the needle holders are preferably tapered from the long sides to the short sides to assist the needle holders in being retracted into the tubular body 12, as will be described in more detail later.

A suture 94, armed at its ends with needles 56, 58, is carried by the device, as seen best in Figure 4. The needles 56, 58 are mounted in the needle carrying grooves 82 and 84 in the needle carriers 70, 72, and the suture 94 is loaded into a suture receiving tube 98 that extends upwardly inside the tubular body 12. As best seen in Figure 6, the needle carriers 70 and 72 and the tube are received in the slots 40, 42 of the actuating member 32. The deep slot 44 is preferably sized to surround the suture holding tube without engaging it, while the shallow slots 40 and 42 are sized and shaped to engage the resilient bars 64, 66 carrying the needle carriers 70 and 72.

When the plunger 22 is fully depressed, the actuating member 32 moves within the tubular body 12 towards the distal end 20 of the trocar wound closure device 10, and pushes the needle carriers 70, 72 to an extended position as shown in Figure 6. When the plunger 22 is withdrawn, the actuating member 32 slides towards the proximal end 14 of the device, and the resilient bars 64, 66 move into the tubular body to withdraw the needle carriers 70, 72 into the body of the trocar wound closure device, as shown in Figure 7, from which the needles and suture are omitted for clarity.

These same two conditions are shown in the section views of the proximal end of the device illustrated at Figures 8 and 9. Figure 8 shows the plunger 22 fully depressed, and Figure 9 shows the plunger 22 fully withdrawn. The housing block 16 is formed from three elements. An inner carrier member 100, preferably cylindrical in configuration and having a through bore therein is mounted within an outer generally rectangular member 104 having a peripheral notch 106 for receiving the top of a cannula and a bore 110 for receiving the inner carrier 100. A flat rectangular cap 112 closes one end of the member, and includes a central bore 114 for receiving the elongated shaft 28 of the plunger 22 therethrough.

The tubular body member 12 is mounted in a slightly enlarged annular recess 116 in the through bore 102 of the inner member 100. The through bore 102 is tapered from a small diameter at its distal end towards a larger diameter at its proximal end. The proximal ends of the resilient bars 64, 66 are attached at radially opposed positions on the inner surface of the tapered portion of the through bore 102. Preferably, the bars 64 and 66 are made from spring steel or a similar resilient material, and are formed so that they naturally assume the position as shown in Figure 9, to retract the needle carriers 70 and 72 to the Figure 7 position, withdrawn into the interior of the distal end 20 of the tubular body 12 when the actuating member 32 is withdrawn to the position shown in Figure 9.

The shallow notches 40, 42 of the actuating element 32 engage the inner surfaces of the bars 64 and 66, as will be recalled from Figure 6. As the plunger 22 is depressed, the actuating element 32 progressively urges the bars 64 and 66 from their natural position to a position adjacent the inner surface of the tubular body 12 as shown in Figures 4, 6 and 8, and at the same time urges the needle carriers 70, 72 to an extended position outside the distal end 20 of the tubular body 12. The upper end of the suture holding tube 98 is also visible in Figures 8 and 9. As shown in Figure 8, a bight formed in the suture 94 may extend slightly beyond the upper end of the suture holder 98, shown shorter than actual size for clarity. As shown in Figure 9, which in this aspect is meant to represent the wound closure device after the suture has been withdrawn through the peritoneal wall, the bight at the center of the suture 94 has disappeared to show that the suture has been withdrawn.

While the trocar wound closure device of this invention may be inserted directly into a trocar wound site, it is conventional in laparoscopic procedures to employ a cannula 120 as shown in Figure 10 during a procedure. The trocar wound closure device of this invention is particularly adapted to be used with the cannula 110 in place at the wound site.

Figure 11 shows the cannula 120 of Figure 10 inserted through a trocar puncture wound. The cannula 110 passes through the skin 124, a fat layer 126 and the fascia 128 into the peritoneal cavity 130.

The invention may be more fully understood by reviewing the manner in which the invention is used to place a suture at a trocar puncture wound site. The suture 94 is armed at each end with a needle 56, 58. As shown in Figures 4 and 8, the center of the suture is stored inside the tube 98 within trocar wound closure device, to keep it from becoming tangled. The needles are mounted in the grooves 82 and 84 of the two needle carriers 70 and 72, as shown in Figure 4, with the sharp ends of the needles pointing towards the proximal end 14 of the trocar wound closure device 10. The suture 94 extends from the distal end of each needle into the body of the trocar wound closure device. The needle carriers 70, 72 are retracted inside the tubular body 12 of the trocar wound closure device to the position shown in Figure 7, so that the outside diameter of the loaded trocar wound closure device is less than the inside diameter of the trocar puncture wound or of the cannula 120.

The trocar wound closure device 10, with the needle carriers retracted, is inserted through the cannula into the peritoneal cavity 130, with the plunger 22 retracted as shown in Figure 11. The plunger 22 is depressed to extend the needle carriers 70, 72 and position the needles 56 and 58 adjacent to but outside the edges of the puncture wound, and in position to puncture the fascia 128 adjacent the wound site as shown in Figure 12. The trocar wound closure device 10 and the cannula 120 are simultaneously pulled outwardly with respect to the cavity 130, and the needles pass through the fascia 128 as shown in Figure 13. The skin 124 and intermediate layer of fat 126 are grasped and retracted from the trocar wound site. The needles 56, 58 are located and pulled only through the fascia and muscle layers, with conventional needle drivers or the like as shown in Figure 14. As the needles and then the suture are pulled through the fascia, the remainder of the suture is controllably released from the suture tube 98 as in Fig. 14. The plunger 22 is first withdrawn to the extended position, prior to withdrwing the trccar wound closure device 10 from the cannula 120. The trocar wound closure device is then retracted from the cannula. The taper of the needle carriers will cause retraction of the needle carriers into the body of the trocar wound closure device as the device is withdraw as shown in Figure 15. After the cannula 120 is withdrawn from the wound a surgical knot is placed over the fascial wound site, thereby closing the wound through the fascia, beneath the fat and skin. The ends of suture are cut. The outer layers of skin and fat are positioned anatomically over the fascial closure, and closed or covered in a conventional manner.

An alternative embodiment of the trocar wound closure device of this invention is shown in Figures 16, 17, 18, and 19. Only the distal end of the device is illustrated in the figures, it being understood that any conventional actuating means as will occur to those skilled in the art may be used. The device shown in these figures includes four needle carriers 202, 204, 206, and 208 rather than two, and is especially suitable for closing large trocar wounds. Four needles 210, 212, 214 and 216 are mounted at the ends of rotatable tubes 220, 222, 224, and 226, carried within the tubular body member 230 of the trocar wound closure device 200. The suture(s) can be stored within one or more of these tubes. The needle holders are elongated and generally oblong in configuration, and each includes an enlarged recess 232, 234, 236, 238 for receiving the end of its respective rotatable tube, and a small recess and slot 240, 242, 244, 246 for receiving a suture armed with a needle. The peripheries of the needle holders are configured so that when rotated to a retracted position as shown in Figure 18, they will fit within the inside of the tubular body 230. At their upper ends, the rotatable tubes are provided with cam shaped actuating mem-

bers 250, 252, 256 and 258, to allow the rods to be rotated to move the needle holders from an extended position, as shown in Figure 17, to a retracted position as shown in Figure 18. Preferably, at least one spacer 260 as shown in Figure 19 is provided within the tubular body member for maintaining the rotatable rods in a spaced apart configuration, while allowing them to turn freely within apertures 262, 264, 266 and 268 in the spacer.

While the invention has been shown and described in connection with a presently preferred embodiment thereof, those skilled in the art will recognize that certain modifications and changes may be made therein without departing from the true scope of the invention.

Claims

1. A trocar wound closure device (10, 200) comprising:

   a) an elongated body (12, 230);
   b) first and second retractable needle holders (70, 72, 202, 204, 206, 208) for holding first and second respective needles (56, 58, 210, 202, 204, 206) therein said needle holders disposed at a distal end (20) of said elongated body and movable between a retracted position such that said needles are maintained substantially within an exterior surface of said elongated body and an extended position wherein said needles are deployed substantially outside said exterior surface substantially parallel to a longitudinal axis of said elongated body;
   c) an open distal end in the elongated body (12), to receive a suture within the elongate body; and
   d) actuator means (32) disposed at a proximal end of the body, for moving the needle holders from the retracted position to the extended position so that the needle holders can be retracted to allow the device to be inserted through a trocar wound, and extended to position the needles adjacent the wound to allow the wound to be closed.

2. A device as claimed in claim 1 wherein said first and second needle holders are configured and dimensioned to deploy first and second needles in a plane substantially parallel to an axial plane of said elongated body when said needle holders are in said extended position.

3. A device as claimed in claim 1 or 2 in which the elongated body comprises a tubular body.

4. A device as claimed in claim 1, 2 or 3 in which the elongated body has openings disposed near said distal end and the retractable needle holders are movable from a retracted position within the body to an extended position outside the body and, extending through the openings.

5. A device as claimed in any one of the preceding claims, in which the retractable needle holders comprise tapered edges which facilitate retraction of the needle holders upon withdrawal of the trocar wound closure device through a cannula.

6. A device as claimed in any one of the preceding claims comprising a tube (98) within the elongated body for carrying a suture.

7. A device as claimed in claim 6 in which the means within the elongated body for carrying a suture includes means for drawing a suture into the tube.

8. A device as claimed in any one of the preceding claims in which the actuator means (32) comprises a plunger (32) movable from an extended position to a depressed position for moving the retractable needle holders from the retracted position to the extended position, respectively.

9. A device as claimed in any one of the preceding claims in which the first and second retractable needle holders comprise first and second needle carriers, slidably mounted in the elongated body, and having needle holding slots (82, 84) oriented generally parallel to a longitudinal axis of the, body, and in which the actuator means comprises means (32) extending through the body for engaging the needle carriers and causing the needle holding slots to move to an extended position.

10. A device as claimed in any one of the preceding claims in which the actuator means comprises a generally cylindrical actuator (32) disposed within the elongated body.

11. A device according to any one of the preceding claims wherein said first and second respective needles are oriented in said needle holders such that pointed ends of said needles are directed generally towards said proximal end of said device.

Patentansprüche

1. Trokarwundenverschlussvorrichtung (10, 200), umfassend:

   a) einen langgestreckten Körper (12, 230);
   b) einen ersten und zweiten einziehbaren Nadelhalter (70, 72, 202, 204, 206, 208), um eine jeweilige erste und zweite Nadel (56, 58, 210, 212, 214, 216) darin zu halten, wobei die besagten Nadelhalter an einem distalen Ende (20) des besagten langgestreckten Körpers angeordnet und zwischen einer eingezogenen

Position, derart, dass die besagten Nadeln im Wesentlichen innerhalb einer äußeren Oberfläche des besagten langgestreckten Körpers gehalten werden, und einer ausgefahrenen Position beweglich sind, in der die besagten Nadeln im Wesentlichen außerhalb der besagten äußeren Oberfläche im Wesentlichen parallel zu einer Längsachse des besagten langgestreckten Körpers ausgebracht sind;

c) ein offenes distales Ende im langgestreckten Körper (12), um einen Faden innerhalb des langgestreckten Körpers aufzunehmen; und

d) eine Betätigungseinrichtung (32), die an einem proximalen Ende des Körpers angeordnet ist, um die Nadelhalter aus der eingezogenen Position in die ausgefahrene Position zu bewegen, so dass die Nadelhalter eingezogen werden können, um ein Einführen der Vorrichtung durch eine Trokarwunde zu ermöglichen, und ausgefahren, um die Nadeln benachbart zur Wunde anzuordnen, um ein Verschließen der Wunde zu ermöglichen.

2. Vorrichtung nach Anspruch 1, dadurch gekennzeichnet, dass der besagte erste und zweite Nadelhalter ausgebildet und bemessen sind, um die erste und zweite Nadel in einer zu einer axialen Ebene des besagten langgestreckten Körpers im Wesentlichen parallelen Ebene auszubringen, wenn sich die besagten Nadelhalter in der besagten ausgefahrenen Position befinden.

3. Vorrichtung nach Anspruch 1 oder 2, dadurch gekennzeichnet, dass der langgestreckte Körper einen röhrenförmigen Körper umfasst.

4. Vorrichtung nach Anspruch 1, 2 oder 3, dadurch gekennzeichnet, dass der langgestreckte Körper nahe dem besagten distalen Ende angeordnete Öffnungen aufweist, und die einziehbaren Nadelhalter aus einer eingezogenen Position innerhalb des Körpers in eine ausgefahrene Position außerhalb des Körpers und durch die Öffnungen verlaufend beweglich sind.

5. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass die einziehbaren Nadelhalter schräge Ränder umfassen, die das Einziehen der Nadelhalter beim Zurückziehen der Trokarwundenverschlussvorrichtung durch eine Kanüle erleichtern.

6. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass sie innerhalb des langgestreckten Körpers ein Rohr (98) zum Tragen eines Fadens umfasst.

7. Vorrichtung nach Anspruch 6, dadurch gekennzeichnet, dass die Einrichtung innerhalb des langgestreckten Körpers zum Tragen eines Fadens Einrichtungen zum Einziehen eines Fadens in das Rohr einschließt.

8. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass die Betätigungseinrichtung (32) einen Plungerkolben (32) umfasst, der aus einer ausgezogenen Position in eine niedergedrückte Position beweglich ist, um die einziehbaren Nadelhalter jeweils aus der eingezogenen Position in die ausgefahrene Position zu bewegen.

9. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass der erste und zweite einziehbare Nadelhalter einen ersten und zweiten Nadelträger umfasst, die verschiebbar im langgestreckten Körper angebracht sind und allgemein parallel zu einer Längsachse des Körpers ausgerichteten Nadelaufnahmeschlitze (82, 84) aufweisen, und dass die Betätigungseinrichtung eine Einrichtung (32) umfasst, die sich durch den Körper erstreckt, um mit den Nadelträgern in Eingriff zu treten und zu bewirken, dass sich die Nadelaufnahmeschlitze in eine ausgefahrene Position bewegen.

10. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass die Betätigungseinrichtung ein innerhalb des langgestreckten Körpers angeordnetes, allgemein zylindrisches Betätigungselement (32) umfasst.

11. Vorrichtung nach einem beliebigen der vorangehenden Ansprüche, dadurch gekennzeichnet, dass die besagte erste und zweite jeweilige Nadel derart in den besagten Nadelhaltern ausgerichtet sind, dass spitze Enden der besagten Nadeln allgemein in Richtung des besagten proximalen Endes der besagten Vorrichtung gerichtet sind.

Revendications

1. Dispositif de fermeture d'une plaie par trocart (10, 200) comprenant :

a) un corps allongé (12, 230);

b) des premier et second supports rétractables d'aiguille (70, 72, 202, 204, 206, 208) pour maintenir des première et seconde aiguilles respectives (56, 58, 210, 202, 204, 206), lesdits supports d'aiguille étant disposés à une extrémité distale (20) dudit corps allongé et mobiles entre une position retirée, telle que lesdites aiguilles soient maintenues sensiblement dans une surface extérieure dudit corps allongé, et une position étendue, où lesdites aiguilles sont déployées sensiblement en dehors de ladite surface extérieure, sensible-

ment parallèlement à un axe longitudinal dudit corps allongé;

c) une extrémité distale ouverte dans le corps allongé ; et

d) un moyen d'actionnement (32) disposé à une extrémité proximale du corps, pour déplacer les supports d'aiguille de la position retirée à la position étendue, de manière que les supports d'aiguille puissent être retirés pour permettre au dispositif d'être inséré à travers une plaie par trocart et étendus pour positionner les aiguilles à proximité de la plaie pour permettre à la plaie de se refermer.

2. Dispositif selon la revendication 1, où lesdits premier et second supports d'aiguille sont configurés et dimensionnés pour déployer les première et seconde aiguilles dans un plan sensiblement parallèle à un plan axial dudit corps allongé quand lesdits supports d'aiguille sont à leur position étendue.

3. Dispositif selon la revendication 1 ou 2, dans lequel le corps allongé comprend un corps tubulaire.

4. Dispositif selon la revendication 1, 2 ou 3, dans lequel le corps allongé a des ouvertures disposées à proximité de ladite extrémité distale et les supports rétractables d'aiguille sont mobiles depuis une position retirée dans le corps à une position étendue en dehors du corps et s'étendant à travers les ouvertures.

5. Dispositif selon lune quelconque des revendications précédentes, dans lequel les supports rétractables d'aiguille comprennent des bords effilés qui facilitent le retrait des supports d'aiguille lors du retrait du dispositif de fermeture d'une plaie par trocart à travers une canule.

6. Dispositif selon l'une quelconque des revendications précédentes, comprenant un tube (98) dans le corps allongé pour porter une suture.

7. Dispositif selon la revendication 6, dans lequel le moyen dans le corps allongé pour porter une suture comporte un moyen pour tirer une suture dans le tube.

8. Dispositif selon l'une quelconque des revendications précédentes, où le moyen d'actionnement (32) comprend un piston (32), mobile d'une position étendue à une position abaissée, pour déplacer les supports rétractables d'aiguille de la position retirée à la position étendue, respectivement.

9. Dispositif selon l'une quelconque des revendications précédentes, dans lequel les premier et second supports rétractables d'aiguille comprennent des premier et second porteurs d'aiguille, montés coulissants dans le corps allongé et ayant des fentes de maintien d'aiguille (82, 84) orientées, généralement, parallèlement à un axe longitudinal du corps et où le moyen d'actionnement comprend un moyen (32) s'étendant à travers le corps pour engager les porteurs d'aiguille et forcer les fentes de soutien d'aiguille à se déplacer à une position étendue.

10. Dispositif selon l'une quelconque des revendications précédentes, dans lequel le moyen d'actionnement comprend un organe d'actionnement généralement cylindrique (32) disposé dans le corps allongé.

11. Dispositif selon l'une quelconque des revendications précédentes, où lesdites première et seconde aiguilles respectives sont orientées dans lesdits supports d'aiguille, de manière que les extrémités pointues desdites aiguilles soient dirigées généralement vers ladite extrémité proximale dudit dispositif.

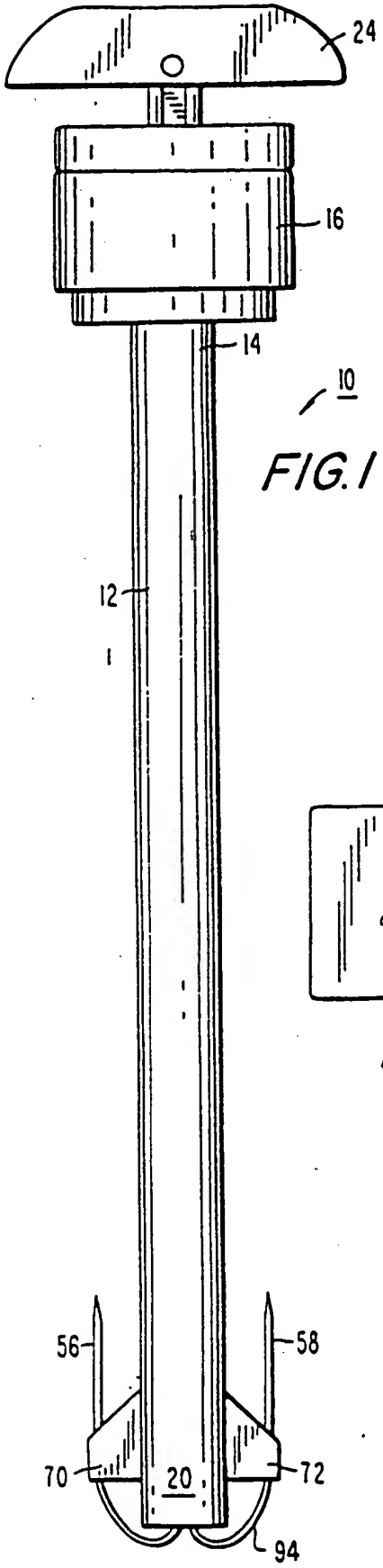


FIG. 1

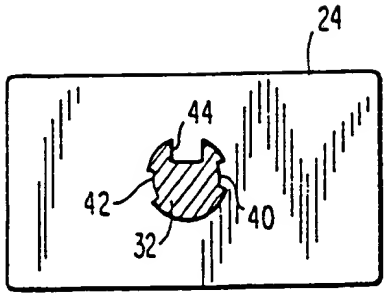


FIG. 3

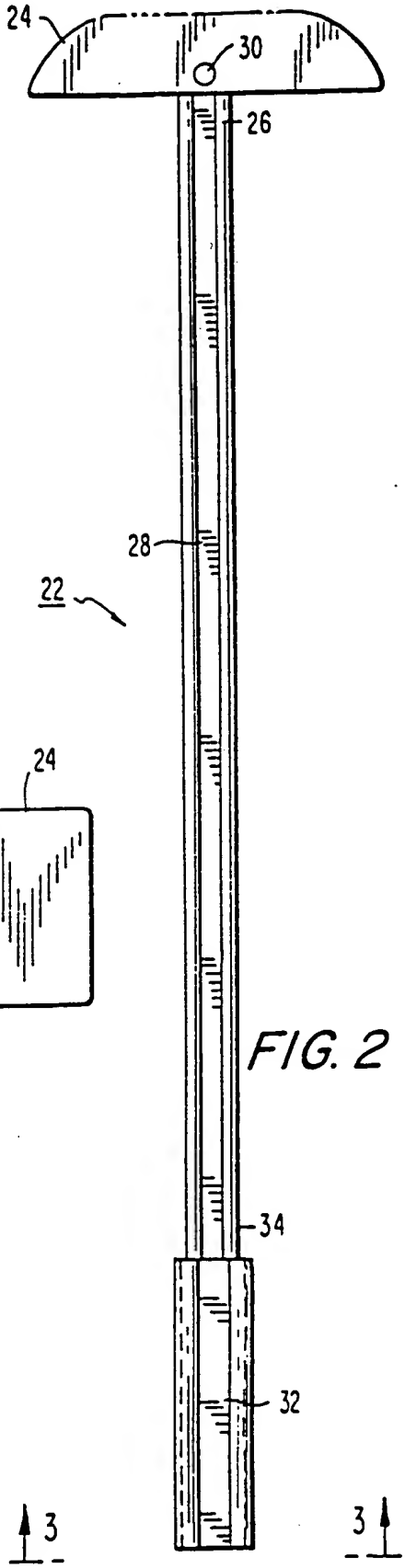


FIG. 2

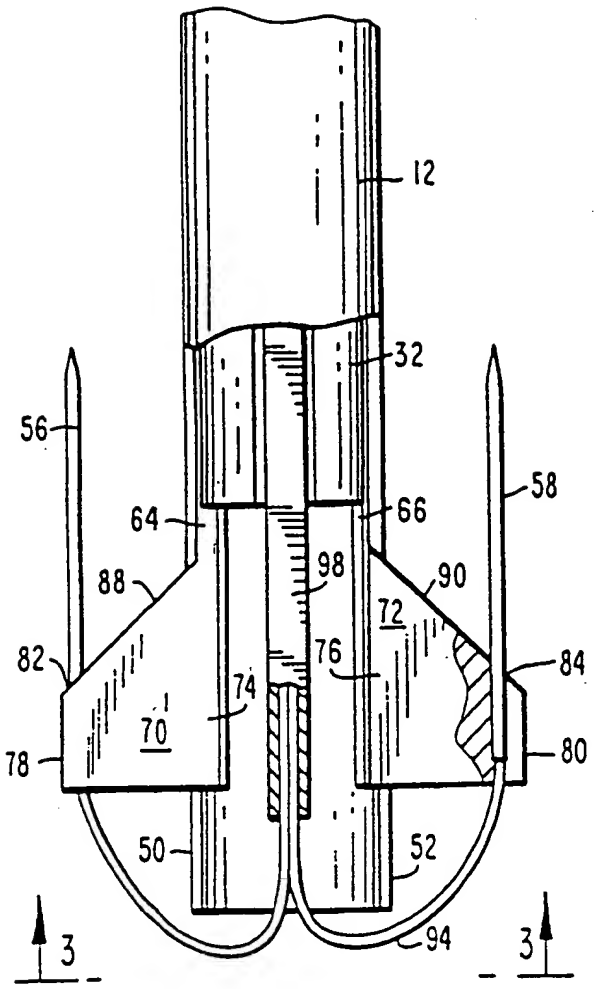


FIG. 4

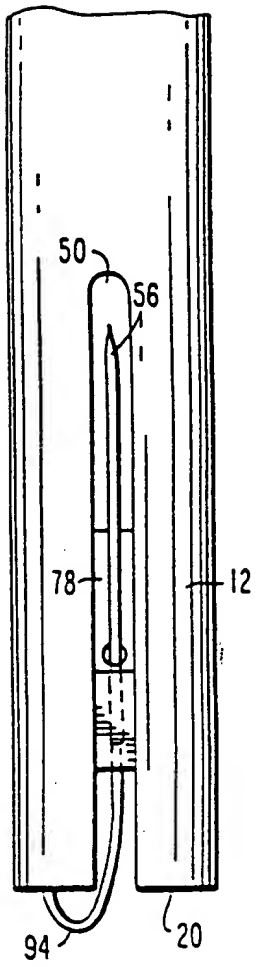


FIG. 5

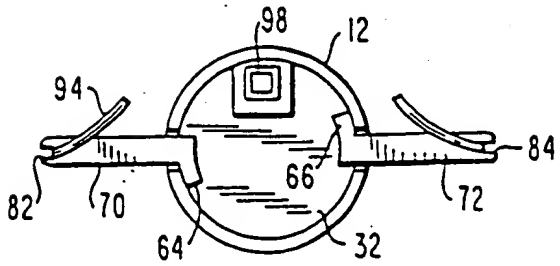


FIG. 6

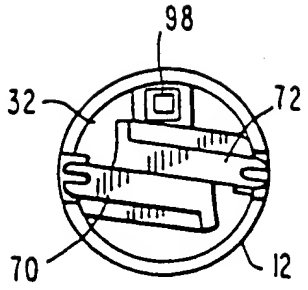
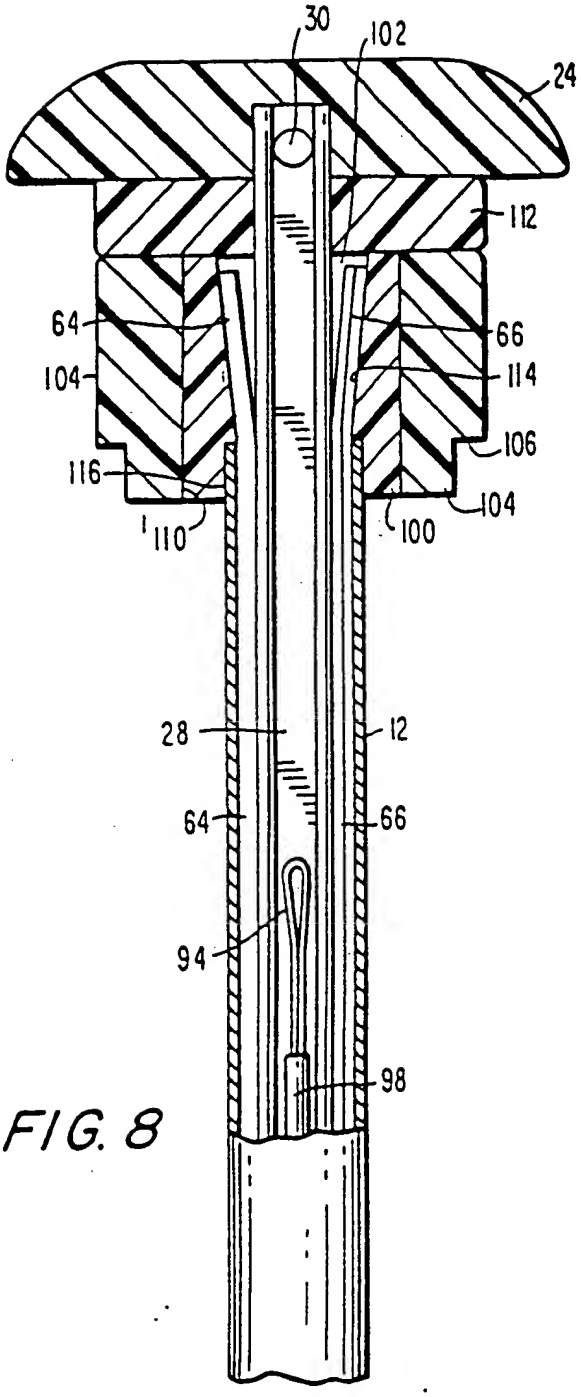


FIG. 7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98

FIG. 8

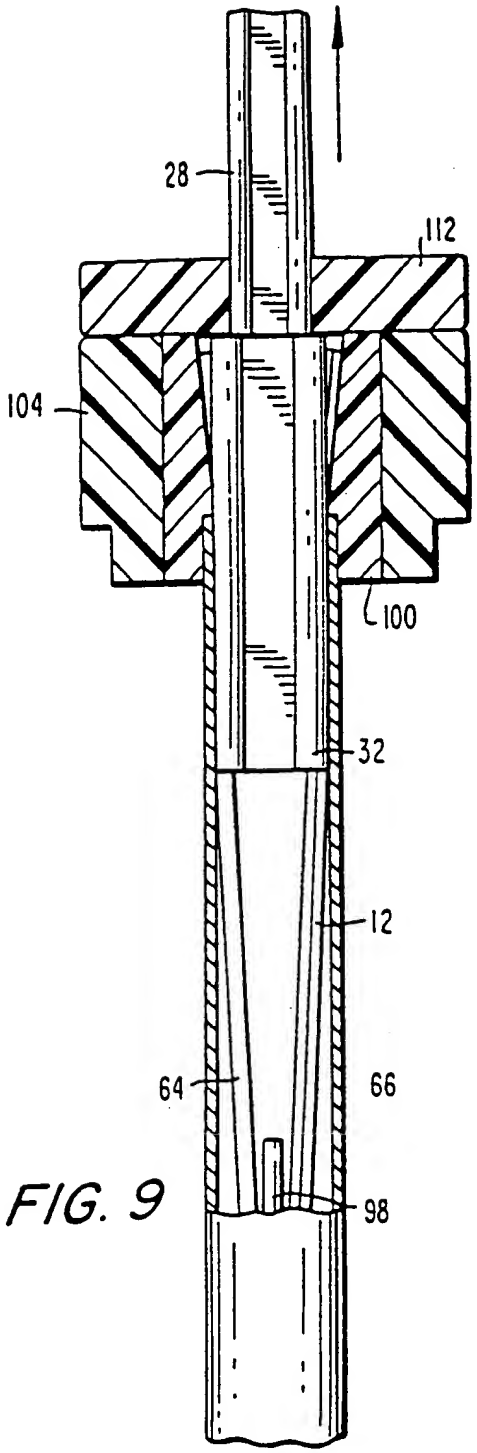
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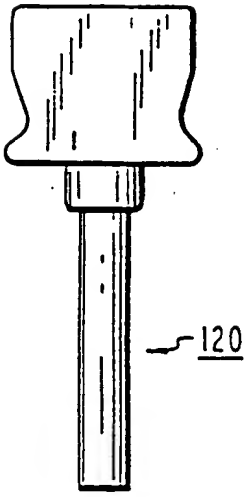
FIG. 9

FIG. 10



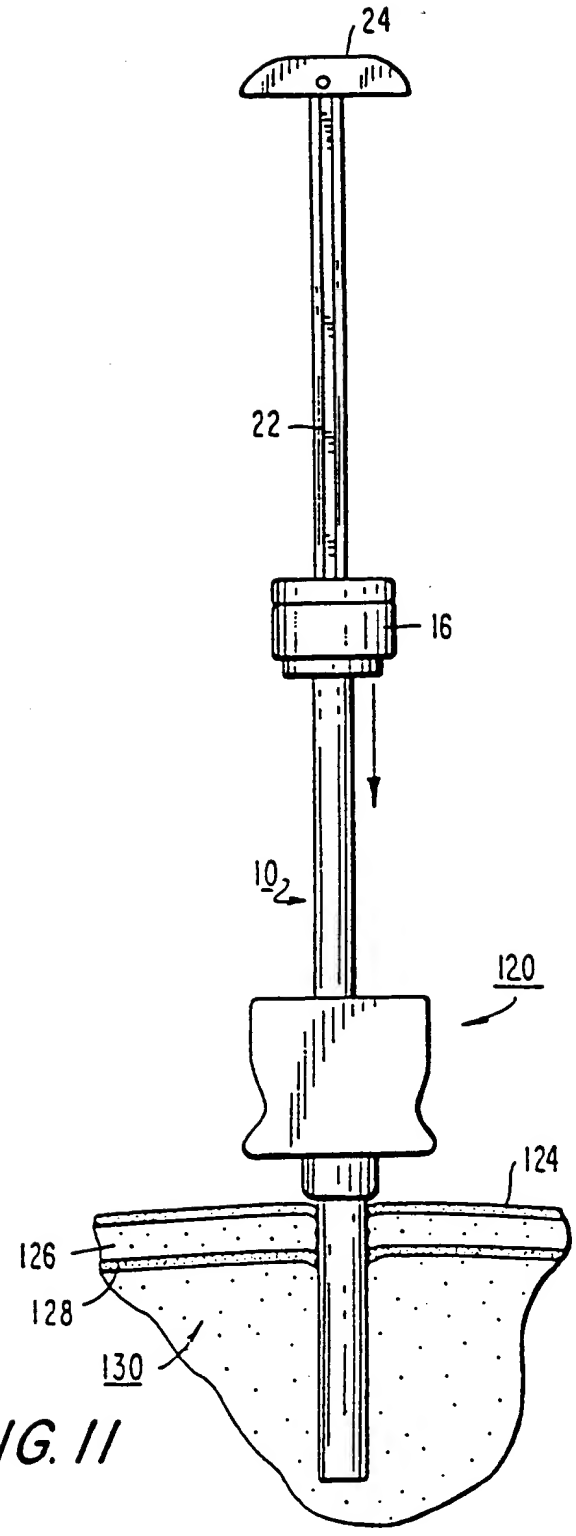


FIG. 11

FIG. 12

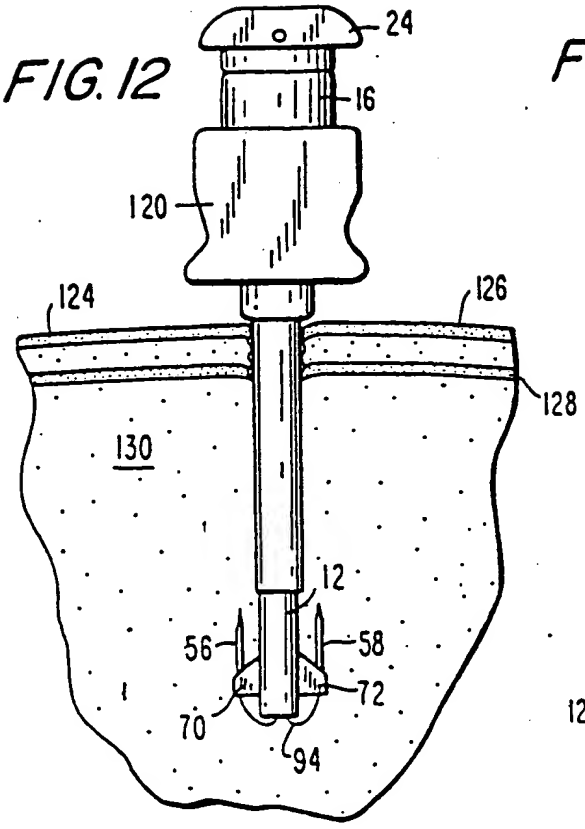


FIG. 13

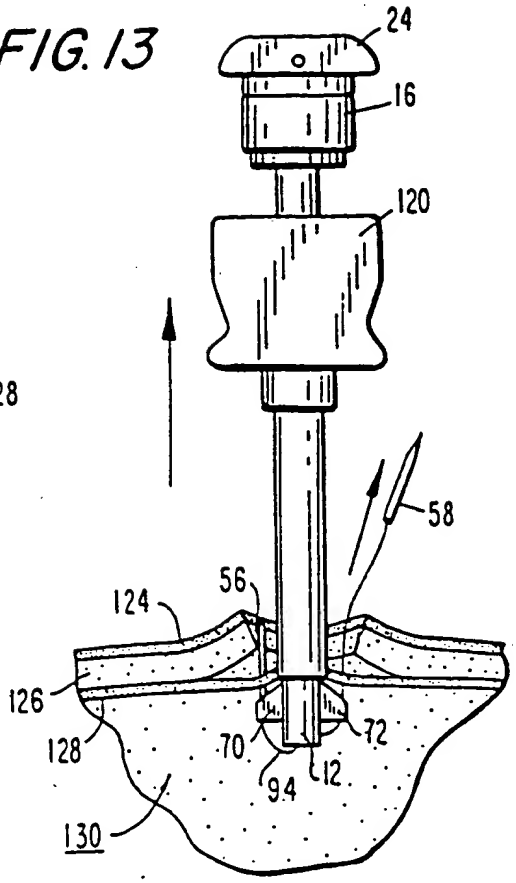


FIG. 14

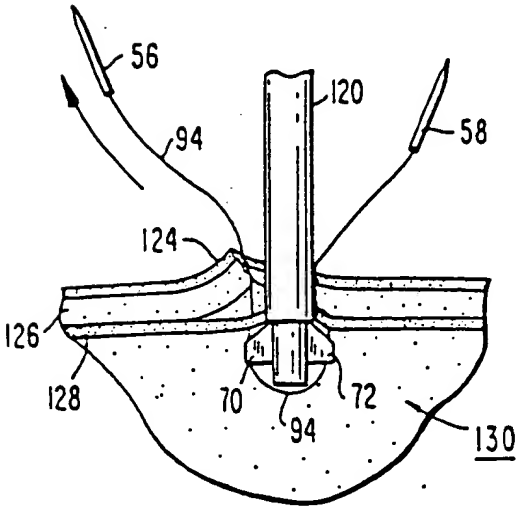
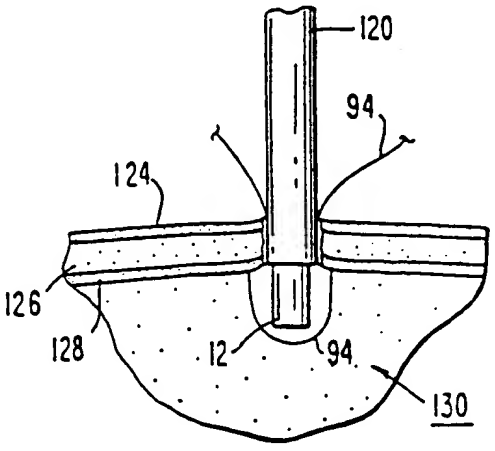


FIG. 15



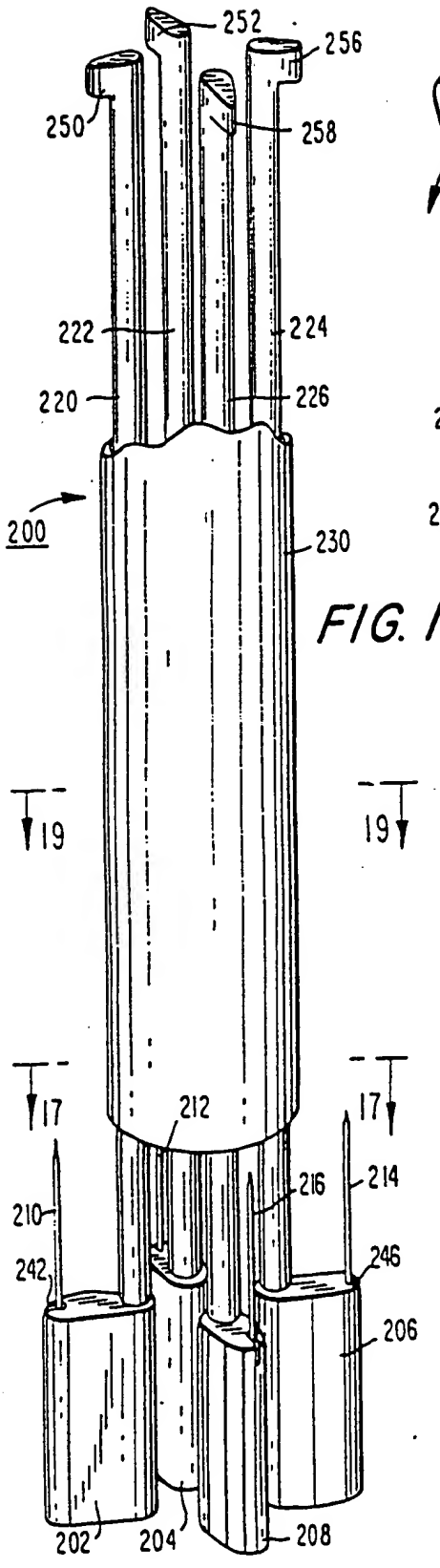


FIG. 16

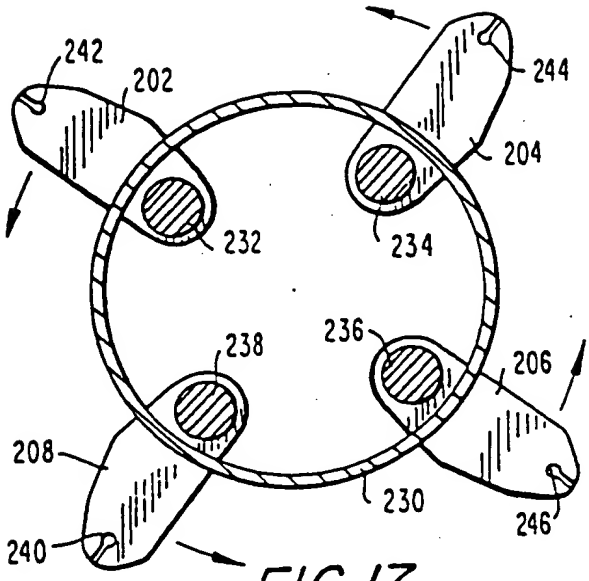


FIG. 17

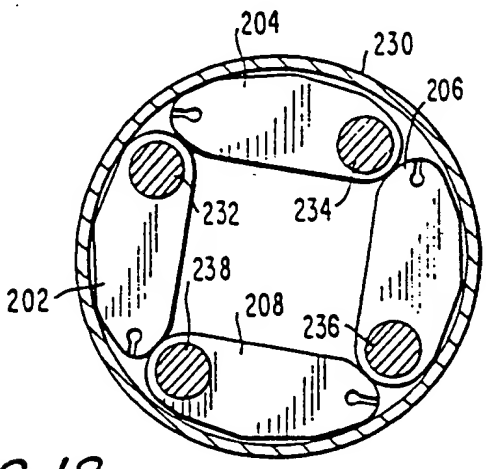


FIG. 18

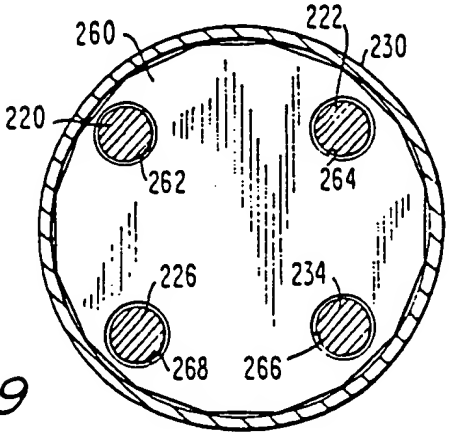


FIG. 19

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Urkunde Certificate Certificat

Es wird hiermit bescheinigt, daß für die in der beigefügten Patentschrift beschriebene Erfindung ein europäisches Patent für die in der Patentschrift bezeichneten Vertragsstaaten erteilt worden ist.

It is hereby certified that a European patent has been granted in respect of the invention described in the annexed patent specification for the Contracting States designated in the specification.

Il est certifié qu'un brevet européen a été délivré pour l'invention décrite dans le fascicule de brevet ci-joint, pour les Etats contractants désignés dans le fascicule de brevet.

Europäisches Patent Nr. **European Patent No.** **Brevet européen n°**

0568098

Patentinhaber **Proprietor of the Patent** **Titulaire du brevet**

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München, den
Munich,
Fait à Munich, le 15.10.97

Ingo Kober

Präsident des Europäischen Patentamts
President of the European Patent Office
Président de l'Office européen des brevets